EXHIBIT 4



Millennium Laboratories

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TEST PROTOCOL - FOR USE WITH POINT OF CARE URINE TEST DEVICES

PLEASE COMPLETE, SIGN	AND FAX	го с	USTOMER	SERVICE AT	858-217-0331
PRACTICE INFORMATIO				SALES RE	
					
				DATE:	
A CONFIRMATION OF NEGATIVE POINT-OF-CARE TEST RESULTS			C ADDITIONAL TESTING NOT INCLUDED ON POINT-OF-CARE TEST		
Millennium Laboratories will confirm and quantify all <u>positive</u> Point-Of-Care results and will confirm negative results as indicated below.		Test SELECT WHICH DRUGS NOT ON THE POINT-OF-CARE TEST YOU WOULD LIKE TESTED AND/OR QUANTIFIED BY LC-MS/MS.			
SELECT WHICH NEGATIVE POINT-OF-CARE RESULTS YOU WOULD LIKE TO HAVE CONFIRMED AND QUANTIFIED BY LC-MS/MS.	ALL		ORDER C	ORDER SCREEN AND C QUANTIFY BY LC-N	
AMPHETAMINES				1V/A	ETHYL GLUCURONIDE
BARBITURATES*					
BENZODIAZEPINES					HEROIN
COCAINE					FENTANYL (DURAGESIC®)
MDMA				닏	CARISOPRODOL (SOMA®)
METHAMPHETAMINE				ᆜ	TRAMADOL (ULTRACET®)
OPIATES					MEPERIDINE (DEMEROL®)
PHENCYCLIDINE				N/A	TAPENTADOL** (NUCYNTA*)
TRICYCLIC ANTIDEPRESSANTS*					and the Transact but to NES/BAS confu
CTHC (MARIJUANA METABOLITE)				by Enzymatic Assay EN VALIDITY T	Only ** Tested by LC-MS/MS only
TO THE PROPERTY OF THE PROPERTY OF CARE THE					perform specimen validity testing (creatinine,
THESE DRUGS MAY OR MAY NOT BE ON YOUR POINT-OF-CARE TEST. PLEASE INDICATE IF YOU WOULD LIKE TO HAVE THESE QUANTIFIED BY LC-MS/M		MS.	pH, spec	cific gravity and o	kidants) unless otherwise indicated here.
ALL			E SPECIAL	INSTRUCTION	S
BUPRENORPHINE (SUBOXONE®)					
METHADONE (METHADOSE®)				·	
OXYCODONE					
PROPOXYPHENE (DARVON®)					
AU I HORIZAI I ON					
1 authorize wineturing transfer as follows:					
individual patient requisition form. 2. Confirm and quantify negative Point-Of-Care Test results as indicated above in Section A, and on the individual patient requisition form. 3. Confirm and quantify all positive Additional Test results pursuant to Section C above. 4. Perform any additional testing (including confirmation of negatives) requested in Sections B and C, and on the individual patient requisition form. 5. Perform specimen validity testing (unless otherwise indicated in Section D).					
		Physician's NPI#: Date://			
		şıgn			
		Physician's Name:			
6. Perform any 'Special Instructions' as specified in Section E.					
I acknowledge that the test panels defined on this Test Protocol are not AMA-approved panels, but nonetheless may be deemed medically necessary because of the clinician's legal and regulatory obligation to take reasonable steps to prevent abuse and diversion of controlled medications. I understand that I may order any of the above-stated tests separately or in necessary combination consistent with the patient's individual medical needs.		Signature: Date:// Physician's Name:			
		Physician's NPI#:			
I understand this Test Pretocol may be modified in writing at any time; otherwise, this Test Protocol will remain in effect for one year from date of signific UMENT 1-4		-			
Test Protocol will remain in effect for one year from date of significant III 1-4		7.GH	GUIGOTZAT	rr caña	2 of 2 Page D #298 / /